



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<b>(54) Title:</b> AQUEOUS AMINOPLAST-CURABLE FILM-FORMING COMPOSITIONS PROVIDING FILMS HAVING RESISTANCE TO ACID ETCHING			
<b>(57) Abstract</b>			
<p>An aqueous aminoplast-curable film-forming composition is disclosed. The aqueous film-forming composition is a cross-linkable composition comprising (1) a material containing a plurality of carbamate and/or urea functional groups and (2) an aminoplast crosslinking agent. The aqueous composition provides a coating with improved acid etch resistance, making the compositions particularly useful as automotive clear coats.</p>			

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AQUEOUS AMINOPLAST CURABLE FILM-FORMING COMPOSITIONS  
PROVIDING FILMS HAVING RESISTANCE TO ACID ETCHING

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Field of the Invention

The present invention relates to aqueous-based, aminoplast curable film-forming compositions, and in particular to aqueous-based aminoplast curable compositions exhibiting superior acid etch 10 resistance.

Background of the Invention

Aminoplast-cured coating systems are well known and provide many excellent coating properties. However, it is widely recognized 15 that such coatings, particularly clear coats, have poor resistance to etching by acid. Conventional coating systems that contain hydroxyl functional film-forming resins and aminoplast crosslinking agents rely on a cure mechanism wherein hydroxyl groups on the resin react with the aminoplast to form ether linkages. See, for example, 20 European Patent Application 0 257 848. Although not intending to be bound by any theory, it is believed that such ether linkages are vulnerable to acid attack and hence yield coatings with poor acid etch resistance.

Because many geographic areas encounter acidic 25 precipitation, acid resistance in coatings is becoming an increasingly desirable property, particularly for automotive coatings. Hydroxyl-aminoplast coating systems of the prior art are not highly effective for providing protection against etching caused by acid rain.

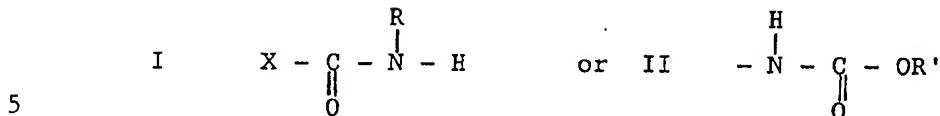
30 It is desirable, therefore, to provide a coating system which avoids the problems of the prior art by demonstrating improved acid etch resistance properties.

Summary of the Invention

35 In accordance with the present invention, an aqueous-based, curable film-forming composition is provided, derived from (1) a

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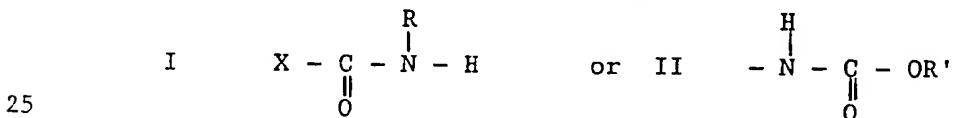
material containing a plurality of terminal or pendant groups of the structure:



where X is  $-N$  or  $-O$  and R is H or alkyl of 1 to 18 carbon atoms or R is bonded to X and forms part of a 5 or 6 membered ring and R' is alkyl of 1 to 18 carbon atoms; and (2) an aminoplast crosslinking agent containing methyol and/or methyol ether groups. Prior to 10 crosslinking, the film-forming composition comprising the material of (1) and (2) has a calculated hydroxyl value less than 50 based on solid weight of the clear film-forming composition, excluding any hydroxyl functionality which may be associated with N-methyol groups. The crosslinked coating has a substantial number of urethane 15 and/or urea crosslinks that arise from reaction of the terminal or pendant groups of structure I or II with the aminoplast, thereby providing a high level of acid etch resistance.

#### Detailed Description

20 The aqueous-based film-forming composition is a crosslinkable composition comprising (1) a material containing a plurality of pendant or terminal groups of the structure:



where X is  $-N$  or  $-O$  and R is H or alkyl of 1 to 18, preferably 1 to 6 carbon atoms or R is bonded to X and forms part of a five- or six-membered ring and R' is alkyl of 1 to 18, preferably 1 to 6 carbon atoms; and (2) an aminoplast crosslinking agent containing 30 methyol and/or methyol ether groups. The material of (1) has on average at least two pendant or terminal groups of the structure I and/or II, preferably structure I, per molecule. Preferably X =  $-O$ . The material of (1) may be an acrylic polymer, a polyester polymer or oligomer, a polyurethane polymer or oligomer, or a blend 35 of two or more of these materials. Acrylic polymers are preferred. Prior to crosslinking, the film-forming composition of (1) and (2)

has a theoretical hydroxyl value of less than 50, preferably less than 25, and more preferably 0, based on solid weight of the film-forming composition, excluding any hydroxyl functionality associated with N-methylol groups such as those in the aminoplast and 5 any hydroxyl functionality which may be associated with N-methylol groups incorporated into the material of (1) such as N-methylol acrylamide groups in the acrylic polymer. By calculated hydroxyl value is meant the calculated value based on the relative amounts of the various ingredients used in making the film-forming composition, 10 rather than the actual hydroxyl value which is measured on the film-forming composition itself by conventional techniques. The resultant crosslinked coating contains a substantial number of urethane or urea crosslinks that arise from reaction of the terminal or pendant groups of structure I or II with the aminoplast, thereby 15 providing a high level of acid etch resistance.

The acrylic materials are copolymers of one or more alkyl esters of acrylic acid or methacrylic acid, and, optionally, one or more other polymerizable ethylenically unsaturated monomers.

Suitable alkyl esters of acrylic or methacrylic acid include methyl 20 methacrylate, ethyl methacrylate, butyl methacrylate, ethyl acrylate, butyl acrylate, and 2-ethylhexyl acrylate. Suitable other polymerizable ethylenically unsaturated monomers include vinyl aromatic compounds such as styrene and vinyl toluene; nitriles such as acrylonitrile and methacrylonitrile; vinyl and vinylidene halides 25 such as vinyl chloride and vinylidene fluoride and vinyl esters such as vinyl acetate. Acrylic and methacrylic acid are also suitable monomers, yielding polymers which can be neutralized and dispersed in water.

Hydroxyl functional monomers such as hydroxyethyl acrylate, 30 hydroxypropyl acrylate, hydroxyethyl methacrylate, and hydroxypropyl methacrylate may be copolymerized with the acrylic monomers to impart hydroxyl functionality to the acrylic material in accordance with the theoretical hydroxyl values mentioned above.

Pendant carbamate functional groups of structure I ( $X = -O$ ) 35 may be incorporated into the acrylic polymer by copolymerizing the acrylic monomers with a carbamate functional vinyl monomer, for

example a carbamate functional alkyl ester of methacrylic acid. These carbamate functional alkyl esters are prepared by reacting, for example, a hydroxyalkyl carbamate, such as the reaction product of ammonia and ethylene carbonate or propylene carbonate, with 5 methacrylic anhydride. Other carbamate functional vinyl monomers are, for instance, the reaction product of hydroxyethyl methacrylate, isophorone diisocyanate, and hydroxypropyl carbamate (yielding structure I), or the reaction product of hydroxypropyl methacrylate, isophorone diisocyanate, and methanol (yielding structure II). Still 10 other carbamate functional vinyl monomers may be used, such as the reaction product of isocyanic acid (HNCO) with a hydroxyl functional acrylic or methacrylic monomer such as hydroxyethyl acrylate, and those described in U.S. Patent 3,479,328. Pendant carbamate groups can also be incorporated into the acrylic polymer by reacting a 15 hydroxyl functional acrylic polymer with a low molecular weight alkyl carbamate such as methyl carbamate. Reference is made to Japanese Kokai 51-4124. Also, hydroxyl functional acrylic polymers can be reacted with isocyanic acid yielding pendant carbamate groups. Note that the production of isocyanic acid is disclosed in U.S. Patent 20 4,364,913. Likewise, hydroxyl functional acrylic polymers can be reacted with urea to give an acrylic polymer with pendant carbamate groups.

Pendant urea groups of structure I ( $X = -N$ ) may be incorporated into the acrylic polymer by copolymerizing the acrylic 25 monomers with urea functional vinyl monomers such as urea functional alkyl esters of acrylic acid or methacrylic acid. Examples include the condensation product of acrylic acid or methacrylic acid with a hydroxyalkyl ethylene urea such as hydroxyethyl ethylene urea. Other urea functional monomers are, for example, the reaction product of 30 hydroxyethyl methacrylate, isophorone diisocyanate, and hydroxyethyl ethylene urea.

Mixed pendant carbamate and urea groups may also be used.

The acrylic polymer material may be prepared by solution polymerization techniques in the presence of suitable catalysts such 35 as organic peroxides or azo compounds, for example, benzoyl peroxide or N,N-azobis(isobutyronitrile). The polymerization may be carried

out in an organic solution in which the monomers are soluble and solubilized or dispersed into water by techniques conventional in the art. Alternately, the acrylic polymer may be prepared directly in water by aqueous emulsion or dispersion polymerization techniques well known in the art. Acrylic polymers with acid functionality can be neutralized for dispersion in water.

The acrylic material typically has a number average molecular weight of from about 900 to 13,000, preferably from about 1000 to 5000 as determined by gel permeation chromatography using a polystyrene standard, and an equivalent weight of less than 5000, preferably within the range of 140 to 2500, based on equivalents of reactive pendant or terminal carbamate or carbamate and/or urea groups. The equivalent weight is a calculated value based on the relative amounts of the various ingredients used in making the acrylic material and is based on solids of the acrylic material.

Polyesters may also be used in the formulation of the film-forming composition and may be prepared by the polyesterification of a polycarboxylic acid or anhydride thereof with polyols and/or an epoxide. Usually, the polycarboxylic acids and polyols are aliphatic or aromatic dibasic acids and diols.

The polyols which are usually employed in making the polyester include alkylene glycols, such as ethylene glycol, 1,6-hexanediol, neopentyl glycol, 2,2-dimethyl-3-hydroxypropyl-2,2-dimethyl-3-hydroxypropionate and other glycols, such as hydrogenated Bisphenol A, cyclohexanediol, cyclohexanedimethanol, caprolactone-based diols, for example, the reaction product of epsilon-caprolactone and ethylene glycol, hydroxy-alkylated bisphenols, polyether glycols, for example, poly(oxytetramethylene) glycol and the like. Polyols of higher functionality may also be used. Examples include trimethylolpropane, trimethylolethane, pentaerythritol and the like.

The acid component of the polyester consists primarily of monomeric carboxylic acids or anhydrides thereof having 2 to 18 carbon atoms per molecule. Among the acids which are useful are phthalic acid, isophthalic acid, terephthalic acid, tetrahydrophthalic acid, hexahydrophthalic acid, adipic acid, azelaic

acid, sebacic acid, maleic acid, glutaric acid, methyl hexahydrophthalic anhydride, decanoic diacid, dodecanoic diacid and other dicarboxylic acids of various types. The polyester may include minor amounts of monobasic acids such as benzoic acid, stearic acid, 5 acetic acid, and oleic acid. Also, there may be employed higher carboxylic acids such as trimellitic acid and tricarballylic acid. Where acids are referred to above, it is understood that anhydrides thereof which exist may be used in place of the acid. Also, lower alkyl esters of the acids such as dimethyl glutarate and dimethyl 10 terephthalate may be used.

Pendant carbamate functional groups of structure I may be incorporated into the polyester by first forming a hydroxyalkyl carbamate which can be reacted with the polyacids and polyols used in forming the polyester. A polyester oligomer may be prepared by 15 reacting a polycarboxylic acid such as those mentioned above with a hydroxyalkyl carbamate. An example of a hydroxyalkyl carbamate is the reaction product of ammonia and ethylene carbonate or propylene carbonate. The hydroxyalkyl carbamate is condensed with acid functionality on the polyester or polycarboxylic acid, yielding 20 pendant carbamate functionality. Pendant carbamate functional groups of structure I may also be incorporated into the polyester by reacting isocyanic acid or a low molecular weight alkyl carbamate such as methyl carbamate with a hydroxyl functional polyester. Also, pendant carbamate functionality may be incorporated into the polyester 25 by reacting a hydroxy functional polyester with urea.

Pendant urea groups of structure I may be incorporated into the polyester by reacting a hydroxyl functional urea such as a hydroxyalkyl ethylene urea with the polyacids and polyols used in making the polyester. A polyester oligomer can be prepared by 30 reacting a polyacid with a hydroxyl functional urea. Also, isocyanate terminated polyurethane or polyester prepolymers may be reacted with primary amines, aminoalkyl ethylene urea, or hydroxyalkyl ethylene urea to yield materials with pendant urea groups. Preparation of these polymers is known in the art and is 35 described in U.S. Patent No. 3,563,957.

Mixed pendant carbamate and urea groups may also be used in the polyester material.

Polyurethanes can be formed by reacting a polyisocyanate with a polyester having hydroxyl functionality and containing the pendant carbamate and/or urea groups. Alternatively, the polyurethane can be prepared by reacting a polyisocyanate with a 5 polyester polyol and a hydroxyalkyl carbamate or isocyanic acid as separate reactants. Examples of suitable polyisocyanates are aromatic and aliphatic polyisocyanates, with aliphatic being preferred because of better color and durability properties. Examples of suitable aromatic diisocyanates are 4,4'-diphenylmethane diisocyanate, 10 1,3-phenylene diisocyanate, 1,4-phenylene diisocyanate, and toluene diisocyanate. Examples of suitable aliphatic diisocyanates are straight chain aliphatic diisocyanates such as 1,4-tetramethylene diisocyanate and 1,6-hexamethylene diisocyanate. Also, cycloaliphatic diisocyanates can be employed and are preferred 15 because of imparting hardness to the product. Examples include 1,4-cyclohexyl diisocyanate, isophorone diisocyanate, alpha,alpha-xylylene diisocyanate and 4,4'-methylene-bis-(cyclohexyl isocyanate).

The polyester and polyurethane materials can contain other 20 pendant groups such as carboxylic acid groups which can be at least partially neutralized with base to aid in water dispersibility. Also, the use of difunctional reactants such as dimethylol propionic acid is possible in the preparation of either a polyester or polyurethane material, providing pendant polar groups.

25 The polyester or polyurethane materials typically have number average molecular weights of about 300 to 3000, preferably about 900 to 1500 as determined by gel permeation chromatography using a polystyrene standard, and an equivalent weight of from about 140 to 2500 based on equivalents of pendant carbamate and/or urea 30 groups. The equivalent weight is a calculated value based on the relative amounts of the various ingredients used in making the polyester or polyurethane and is based on solids of the material.

Besides polymeric materials, relatively low molecular weight materials containing pendant carbamate functional groups of structure 35 II may be formed by reacting isocyanate terminated monomers or oligomers, such as an isocyanurate of polymeric 1,6-hexamethylene

diisocyanate, with an alcohol. Any suitable aliphatic, cycloaliphatic, aromatic alkyl monoalcohol or phenolic compound may be used, such as, for example, aliphatic alcohols containing from 1 to 18, preferably lower aliphatic alcohols containing from 1 to 6 carbon atoms such as methanol, ethanol, n-butyl alcohol and n-hexanol; cycloaliphatic alcohols such as cyclohexanol; aromatic-alkyl alcohols such as phenyl carbinol and methylphenyl carbinol; phenolic compounds such as phenol itself, and substituted phenols in which the substituents do not adversely affect coating operations. Examples include cresol and nitrophenol.

It is possible to prepare blends of the acrylic, polyester, and polyurethane materials containing pendant or terminal carbamate and/or urea groups described above. It is also possible to prepare blends of the low molecular weight materials containing pendant carbamate and/or urea groups with the polymeric materials containing pendant carbamate and/or urea groups. The weight ratio of low molecular weight materials to polymeric materials may range from 10:90 to 90:10, preferably 10:90 to 40:60.

The aqueous film-forming composition also includes an aminoplast crosslinking agent containing methylol and/or methylox ether groups. Aminoplast condensates are obtained from the reaction of formaldehyde with an amine or amide. The most common amines or amides are melamine, urea, or benzoguanamine, and are preferred. While the aldehyde used is most often formaldehyde, other aldehydes such as acetaldehyde, crotonaldehyde, and benzaldehyde may be used.

The aminoplast contains methylol groups and preferably at least a portion of these groups are etherified with an alcohol to modify the cure response. Any monohydric alcohol may be employed for this purpose including methanol, ethanol, butanol, and hexanol.

Preferably, the aminoplasts which are used are melamine-, urea-, or benzoguanamine-formaldehyde condensates etherified with an alcohol containing from 1 to 6 carbon atoms. The aminoplast is present in amounts of about 1 to 80, preferably 10 to 50 percent by weight based on weight of resin solids in the clear film-forming composition. The equivalent ratio of pendant or terminal carbamate and/or urea functional groups of structure I and II above to methylol

or methylol ether groups is 0.5 to 2:1 based on calculated equivalent weights, and being sufficient to form a crosslinked film.

The film-forming composition can be dispersed in water in several ways. As mentioned above, acid-functional materials having 5 terminal or pendant carbamate and/or urea groups may be neutralized with bases such as amines and dissolved or dispersed in water. Also, it is possible to prepare an aqueous dispersion of a blend of acrylic and polyester and/or polyurethane materials with pendant carbamate and/or urea groups in microparticulate form by a high stress 10 technique using a homogenizer. This technique is described in U.S. Patent No. 5,071,904.

The aqueous, liquid film-forming composition will also preferably contain catalysts to accelerate the cure of the aminoplast and carbamate or urea groups. Examples of suitable catalysts are 15 acidic materials and include sulfonic acids or substituted sulfonic acids such as para-toluenesulfonic acid. The catalyst is usually present in an amount of about 0.5 to 5.0 percent by weight, preferably about 1 to 2 percent by weight, based on weight of total resin solids. Optional ingredients such as, for example, 20 plasticizers, flow controllers, anti-oxidants, UV light absorbers and similar additives conventional in the art may be included in the composition. These ingredients are typically present at up to 25% by weight based on total resin solids.

The aqueous-based composition may be applied to a substrate, 25 or in the case of a clear film-forming composition, to a basecoated substrate by any conventional coating technique such as brushing, spraying, dipping or flowing, but spray applications are preferred because of superior gloss. Any of the known spraying techniques may be employed such as compressed air spraying, electrostatic spraying 30 and either manual or automatic methods.

After application of the coating composition, the coated substrate is heated to cure the coating. In the curing operation, solvents and water are driven off and the film-forming material of the coating is crosslinked. The heating or curing operation is 35 usually carried out at a temperature in the range of from 160-350°F (71-177°C) but if needed, lower or higher temperatures may be used as

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necessary to activate crosslinking mechanisms. The thickness of the coating is usually from about 0.5-5, preferably 1.2-3 mils.

The invention will further be described by reference to the following examples. Unless otherwise indicated, all parts are by weight.

#### EXAMPLES

The following examples (Examples A-I) show the preparation of carbamate and/or urea functional materials and corresponding hydroxyl functional materials.

##### Example A

A carbamate functional acrylic monomer was prepared from the following ingredients:

	<u>Ingredient</u>	<u>Weight in Grams</u>
15	isophorone diisocyanate (IPDI)	888.0
	dibutyl tin dilaurate	4.6
	2,6-di-t-butyl methyl phenol	2.6
	butyl methacrylate	282.0
20	hydroxypropyl carbamate	571.2
	hydroxyethyl methacrylate	416.0

A suitable reactor was charged with the first four ingredients and heated to a temperature of 60°C. The hydroxypropyl carbamate was added to the reaction mixture over 2 hours. The reaction mixture was then held at 60°C until the isocyanate equivalent weight became constant. The hydroxyethyl methacrylate was then added over 2 hours, and the reaction held until infrared analysis indicated the absence of isocyanate. The product was diluted with 346.0 g of butyl methacrylate. The final product had a solids content of 75% and had a number average molecular weight of 622 as determined by gel permeation chromatography.

##### Example B

A carbamate functional acrylic polymer dispersed in aqueous medium was prepared from the following ingredients:

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<u>Ingredient</u>	<u>Weight in Grams</u>
n-propanol	350.0
butyl acrylate	202.0
methyl methacrylate	195.2
5 carbamate functional acrylic monomer from Example A	349.9
acrylic acid	25.0
t-dodecyl mercaptan	3.2
t-butyl peroctoate	14.4
10 n-propanol	46.4
dimethyl ethanol amine (DMEA)	23.2
water	700.0

A suitable reactor was charged with the n-propanol and heated to reflux. The next five ingredients were added to the 15 reaction mixture over 3 hours. At the same time, the t-butyl peroctoate and 46.4 g n-propanol were added over 3.5 hours. The reaction was then held at reflux for one hour. The DMEA was added to the reaction mixture at about 95°C, followed by addition of the water. The reaction cooled to room temperature. The final product 20 had a solids content of 35.3% and had a number average molecular weight of 3728 as determined by gel permeation chromatography. The carbamate equivalent weight of the resultant material was approximately 1040.

Example C

25 A carbamate functional acrylic latex was prepared from the following ingredients:

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	<u>Ingredient</u>	<u>Weight in Grams</u>
Feed A:	water	783.4
	ALIPAL CO-436 <sup>1</sup>	15.1
	sodium bicarbonate	1.8
5	Feed B: water	114.8
	ammonium persulfate	5.2
Feed C:	butyl acrylate	277.5
	methyl methacrylate	263.7
10	carbamate functional acrylic monomer from Example A	502.0
	butyl methacrylate	136.9
	acrylic acid	36.4
	t-dodecyl mercaptan	18.2
	water	757.7
15	ALIPAL CO-436	17.4
	DDBSA-DMEA <sup>2</sup>	11.5
Feed D:	diisopropanol amine, 50% in water	67.2
1	Anionic ethoxylated nonyl phenol available from GAF Corporation.	
2	DDBSA-DMEA solution was prepared by dissolving 1 mole dodecyl	
20	benzene sulfonic acid in water containing 1 mole dimethyl ethanolamine.	
25	A suitable reactor was charged with Feed A and heated to 80°C. 25 g of Feed C and then all of Feed B were added to the reaction mixture, and the mixture was held for 20 minutes. The remainder of Feed C was added over 3 hours. The reaction was held at 80°C for two hours, and then cooled to room temperature. After dilution with Feed D, the final product had a solids content of 42.8%	
	and had a number average molecular weight of 12,393 as determined by gel permeation chromatography. The carbamate equivalent weight of	
30	the resultant material was approximately 1140.	

#### Example D

A carbamate functional polyester was prepared from the  
following ingredients:

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<u>Ingredient</u>	<u>Weight in Grams</u>
DOWANOL PM carbamate <sup>2</sup>	332.5
polyester <sup>1</sup>	455
butyl stannoic acid	2.12
5        1 Reaction product of hexahydrophthalic anhydride, ESTERDIOL 204, and 1,6-hexanediol in a 1:1:1 mole ratio.	

2        2 Reaction product of DOWANOL PM and urea, 95% in DOWANOL PM which is the monomethyl ether of propylene glycol and is available from the Dow Chemical Co.

10        The ingredients were charged to a suitable reactor equipped with a nitrogen sparge and Dean-Stark trap and heated to 140-145°C. DOWANOL PM was removed from the reaction under reduced pressure. The reaction mixture was held until DOWANOL PM carbamate was no longer detectable on a gas chromatograph. The resultant reaction mixture 15 was a soft, waxy, opaque material.

#### Example E

A pre-emulsion was prepared by stirring together the following ingredients:

20	<u>Ingredient</u>	<u>Weight in Grams</u>
	carbamate functional polyester of Example D	125.0
	methyl methacrylate	100.0
	butyl acrylate	100.0
25	stearyl methacrylate	25.0
	N-methylo1 acrylamide (48% solution in water)	83.4
	methacrylic acid	10.0
	dodecylbenzenesulfonic acid (70% in water)	14.3
30	N,N-dimethyl ethanol amine	2.5
	IGEPAL CO-897 <sup>1</sup>	7.2
	ferrous ammonium sulfate, 1% in water	2.5
	water	500.0

35        1 Nonionic ethoxylated nonyl phenol available from GAF Corp.

The pre-emulsion was passed though an M110 MICROFLUIDIZER high pressure impingement emulsifier (available from Microfluidics,

Inc.) at 8000 psi to produce a bluish-white emulsion. The emulsion was transferred to a suitable reactor and blanketed with nitrogen. Polymerization was initiated by adding first a mixture of 1.5 g isoascorbic acid and 2.5 g mercaptopropionic acid dissolved in 50.0 g water followed by a solution of 2.19 g hydrogen peroxide (35%) in 25.0 g water added dropwise over 15 minutes. The emulsion exothermed from 26 to 66°C over 14 minutes. Any remaining monomer was then polymerized by adding 0.5 g isoascorbic acid dissolved in 5.0 g water followed by 0.5 g of 35% hydrogen peroxide. An additional exotherm from 56 to 59°C was observed. The pH of the latex was increased to 10 7.0 with 16.45 g of a 1:1 mixture of water and diisopropanolamine. The final product had a solids content of 41.0%.

Example F

15 A urea functional polyester oligomer was prepared from the following ingredients:

	<u>Ingredient</u>	<u>Weight in Grams</u>
	dodecanedioic acid	575.0
	hydroxyethyl ethylene urea	637.74
20	butyl stannoic acid	1.21
	xylene	198.66

The ingredients were charged to a suitable reactor and heated to reflux to remove water through a Dean-Stark trap. The temperature of the reaction mixture was held at reflux until the acid 25 value was less than 5. The reaction mixture was then cooled to 120°C and volatile materials in the reaction mixture were removed under vacuum to a solids content of 98.7%. The reaction mixture was diluted to a final solids content of 65% with an 80:20 weight mixture of propanol:water. The product had a number average molecular weight 30 of 606 and a theoretical urea equivalent weight of approximately 230.

Example G

A carbamate functional acrylic monomer was prepared from the following ingredients:

<u>Ingredient</u>	<u>Weight in Grams</u>
hydroxypropyl carbamate	600.0
2,6-di-t-butyl methyl phenol	3.9
triphenyl phosphite	2.22
5 methacrylic anhydride	810.0
toluene	1200.0
sodium hydroxide (16.7%)	1260.0

A suitable reactor was charged with the first four ingredients and heated to 100°C. The reaction mixture was held at 10 this temperature until the methacrylic anhydride had completely reacted with the hydroxypropyl carbamate, as determined by gas chromatography. The reaction was cooled to room temperature and the toluene and sodium hydroxide were added. After agitating for about 30 minutes, the reaction mixture was transferred to a separatory 15 funnel. The top layer, containing the product in toluene, was collected in a flask and the toluene was removed by vacuum distillation.

#### Example H

20 A carbamate functional acrylic latex was prepared from the following ingredients:

	<u>Ingredient</u>	<u>Weight in Grams</u>
Feed A:	water	450.0
	ALIPAL CO-436	9.3
25	sodium bicarbonate	0.8
Feed B:	water	50.0
	ammonium persulfate	2.2
Feed C:	carbamate functional acrylic monomer of Example G	180.0
30	butyl acrylate	240.0
	methyl methacrylate	120.0
	styrene	60.0
	acrylic acid	16.8
	t-dodecyl mercaptan	9.0
35	water	400.0
	ALIPAL CO-436	18.0
	PGNP-15 <sup>1</sup>	26.0
Feed D:	diisopropanol amine, 50% in water	20.0

<sup>1</sup> Nonionic surfactant prepared by reacting 1 mole of nonyl phenol with 15 moles of glycidol.

A suitable reactor was charged with Feed A and heated to 80°C. 25 g of Feed C and then all of Feed B were added to the reaction mixture, and the mixture was held for 20 minutes. The remainder of Feed C was added over 3 hours. The reaction was held at 80°C for two hours, and then cooled to room temperature. After addition of Feed D, the pH was 7.7. The final product had a solids content of 40.5% and had a number average molecular weight of 5706 as determined by gel permeation chromatography.

#### Example I

A hydroxyl functional acrylic latex was prepared from the following ingredients:

	<u>Ingredient</u>	<u>Weight in Grams</u>
15	Feed A: water	450.0
	ALIPAL CO-436	9.3
	sodium bicarbonate	0.8
20	Feed B: water	50.0
	ammonium persulfate	2.2
25	Feed C: hydroxyethyl acrylate	180.0
	butyl acrylate	240.0
	methyl methacrylate	120.0
	styrene	60.0
	acrylic acid	16.8
30	t-dodecyl mercaptan	9.0
	water	400.0
	ALIPAL CO-436	18.0
	PGNP-15	26.0
Feed D:	diisopropanol amine, 50% in water	20.0

A suitable reactor was charged with Feed A and heated to 80°C. 25 g of Feed C and then all of Feed B were added to the reaction mixture, and the mixture was held for 20 minutes. The remainder of Feed C was added over 3 hours. The reaction was held at 80°C for two hours, and then cooled to room temperature. After addition of Feed D, the pH was 7.84. The final product had a solids

content of 40.2% and had a number average molecular weight of 5123 as determined by gel permeation chromatography, and a hydroxyl value of 22 based on solids content.

5 The following examples (1-6) show the preparation of various clear film-forming compositions prepared with carbamate, urea, or hydroxyl functional materials and aminoplast curing agents. The coating compositions were evaluated in color-plus-clear applications.

10

Example 1

A clear film-forming composition was prepared by mixing together the following ingredients:

<u>Ingredient</u>	<u>Solid weight in grams</u>	<u>Solution weight in grams</u>
TINUVIN 1130	3.5	3.5
15 CYMEL 328 <sup>1</sup>	30.0	34.9
carbamate containing acrylic of Example B	70.0	198.4
phenyl acid phosphate	1.0	5.0
water	---	137.0

20 <sup>1</sup> Waterborne version of CYMEL 327 available from American Cyanamid.

Example 2

A clear film-forming composition was prepared by mixing 25 together the following ingredients:

<u>Ingredient</u>	<u>Solid weight in grams</u>	<u>Solution weight in grams</u>
carbamate containing acrylic of Example C	70.0	162.6
CYMEL 303 <sup>1</sup>	30.0	30.0
30 TINUVIN 1130	3.5	3.5
DDBSA solution <sup>2</sup>	1.0	5.0
FC 430 solution <sup>3</sup>	0.1	2.0
diisopropanol amine solution <sup>4</sup>	---	3.9
n-methyl-2-pyrrolidone	---	5.0
35 isopropanol	---	5.0
water	---	25.0

<sup>1</sup> Hexamethoxymethyl melamine resin available from American Cyanamid.

<sup>2</sup> 20 weight percent solution of dodecylbenzene sulfonic acid neutralized with diisopropanolamine in deionized water.

<sup>3</sup> Nonionic surfactant available from 3M Corporation.

<sup>4</sup> 50 weight percent solution of diisopropanolamine in deionized water.

### Example 3

10 A clear film-forming composition was prepared by mixing together the following ingredients:

	<u>Ingredient</u>	<u>Solid weight in grams</u>	<u>Solution weight in grams</u>
	DDDA/HEEU oligomer of Example F	70.0	116.7
15	CYTEL 328	30.0	34.9
	Phenyl acid phosphate solution	1.0	5.0
	Tego Wet ZFS 453 <sup>1</sup>	0.09	0.36

<sup>1</sup> Nonionic surfactant available from Tego Chemie Service GmbH.

20

### Example 4

A clear film-forming composition was prepared by mixing together the following ingredients:

	<u>Ingredient</u>	<u>Solid weight in grams</u>	<u>Solution weight in grams</u>
25	carbamate functional acrylic and polyester latex of Example E	70.0	170.61
	CYTEL 303	30.0	30.0
	TINUVIN 1130	3.5	3.5
30	DDBSA solution	1.0	5.0
	FC 430 solution	0.1	2.0
	diisopropanol amine solution	--	3.2
	n-methyl-2-pyrrolidone	--	5.0
	isopropanol	--	5.0
35	water	--	58.1

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Example 5

A clear film-forming composition was prepared by mixing together the following ingredients:

<u>Ingredient</u>	<u>Solid weight in grams</u>	<u>Solution weight in grams</u>
5 carbamate functional acrylic latex of Example H	70.0	172.8
CYMEL 303	30.0	30.0
TINUVIN 1130	3.5	3.5
p-TSA solution <sup>1</sup>	1.0	5.0
10 diisopropanol amine solution	--	3.5
isopropanol	--	50.0
water	--	17.3

<sup>1</sup> 20 weight percent solution of para-toluene sulfonic acid neutralized with diisopropanol amine in water.

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Example 6

A clear film-forming composition was prepared by mixing together the following ingredients:

<u>Ingredient</u>	<u>Solid weight in grams</u>	<u>Solution weight in grams</u>
20 hydroxyl functional acrylic latex of Example I	70.0	174.2
CYMEL 303	30.0	30.0
TINUVIN 1130	3.5	3.5
p-TSA solution	1.0	5.0
25 diisopropanol amine solution	--	3.5
isopropanol	--	50.0
water	--	16.3

The film-forming compositions of Examples 1-6 were applied to a pigmented basecoat to form color-plus-clear composite coatings over electrocoated steel substrates. The pigmented basecoat for Examples 1-6 is commercially available from PPG Industries, Inc. and identified as BWB-8555. The basecoat was pigmented black in color. The electrocoat used on the steel is commercially available from PPG Industries, Inc. and is identified as ED-11.

35 The basecoat was spray applied in two coats to electrocoated steel panels at a temperature of about 75°F (24°C) and a relative

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humidity of about 60%. A ninety second flash time was allowed between the two basecoat applications. After the second basecoat application, a prebake time of approximately five minutes was allowed at 250°F (121°C) before the application of the clear coating  
5 composition. The clear coating compositions of Examples 1-6 were each applied to a basecoated panel in two coats with a ninety second flash at 75°F (24°C) allowed between coats. The composite coating was allowed to air flash at 75°F (24°C) for ten to fifteen minutes and to flash at 140°F (60°C) for ten to fifteen minutes before baking  
10 at 285°F (141°C) for 30 minutes to cure both the basecoat and clearcoat. The panels were baked in a horizontal position. The properties of the composite coatings are reported in Table I below.

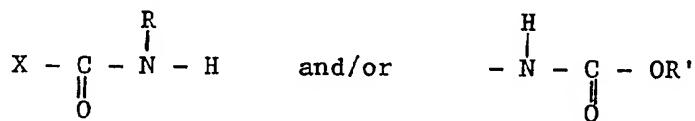
Table I

	<u>Example</u>	<u>Acid etch rating</u>
15	1	3
	2	3
	3	2
	4	5
	5	5
20	6	9

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We claim:

1. An aqueous-based curable liquid film-forming composition comprising (1) a material containing a plurality of 5 terminal or pendant groups of the structure:



where X is -N or -O and R is H or alkyl of 1-18 carbon atoms or R is bonded to X and forms part of a 5 or 6 membered ring and R' is alkyl of 1 to 18 carbon atoms; and (2) an aminoplast crosslinking agent containing methylol and/or methylol ether groups; said film-forming composition being crosslinkable through reaction of said pendant or terminal groups with said methylol and/or methylol ether groups; said film-forming composition being further characterized as having a calculated hydroxyl value less than 50 based on solid weight of said film-forming composition, excluding any hydroxyl functionality associated with N-methylol groups so as to result in a crosslinked coating which has a substantial number of urethane and/or urea crosslinks arising from said reaction of pendant or terminal groups with said methylol and/or methylol ether groups, giving said crosslinked coating a high level of acid etch resistance.

2. The composition of claim 1 in which the material is an acrylic polymer.

3. The composition of claim 2 in which the acrylic polymer is prepared by aqueous emulsion polymerization techniques.

4. The composition of claim 2 in which the acrylic polymer is prepared in organic solvent and inverted into aqueous medium.

5. The composition of claim 4 in which the acrylic polymer contains ionic groups.

35

6. The composition of claim 5 in which the ionic groups are anionic groups.

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7. The composition of claim 2 in which the acrylic polymer has an equivalent weight less than 5000 based on equivalents of said terminal or pendant groups.

5 8. The composition of claim 7 in which the acrylic polymer has an equivalent weight of from about 140 to 2500 based on equivalents of said terminal or pendant groups.

10 9. The composition of claim 1 in which the material is selected from the class consisting of polyesters, polyurethanes, and mixtures thereof.

10. The composition of claim 9 in which the material is prepared in organic solvent and inverted into aqueous medium.

15

11. The composition of claim 10 in which the material contains ionic groups.

12. The composition of claim 11 in which the ionic groups 20 are anionic groups.

13. The composition of claim 9 in which the material has on average at least two of said pendant or terminal groups.

25 14. The composition of claim 13 in which the material has an equivalent weight of from about 140 to 2500 based on equivalents of said terminal or pendant groups.

30 15. The composition of claim 1 in which X is -O and R is selected from the class consisting of H and alkyl containing from 1 to 6 carbon atoms and R' is alkyl containing from 1 to 6 carbon atoms.

35 16. The composition of claim 1 in which the aminoplast is a condensate of melamine with formaldehyde and optionally an alcohol containing from 1 to 6 carbon atoms.

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17. The composition of claim 15 wherein the equivalent ratio of said pendant or terminal groups to methylol or methylo1 ether groups is from 0.5 to 2:1 and is sufficient to form a crosslinked film.

5

18. The composition of claim 1 wherein the material is present in the film-forming composition in amounts of 50 to 90 percent by weight based on weight of resin solids in the film-forming composition.

10

19. The composition of claim 1 wherein the aminoplast is present in the film-forming composition in amounts of 10 to 50 percent based on weight of resin solids in the film-forming composition.

# INTERNATIONAL SEARCH REPORT

Int'l Application No  
PCT/US 93/10173

**A. CLASSIFICATION OF SUBJECT MATTER**  
IPC 5 C08F8/30 C09D201/02

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)  
IPC 5 C08F C09D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO,A,87 00851 (UCC) 12 February 1987 see page 11 see page 15; claims 1-56 ---	1-19
A	US,A,3 984 382 (G. G. PAREKH) 5 October 1976 see column 7, line 41 - column 8, line 32; claims 1-10 ---	1
A	US,A,3 959 201 (PPG INDUSTRIES, INC.) 25 May 1976 see claims 1-19 ---	1
A	WO,A,91 08269 (BOLLIG & KEMPER KG) 13 June 1991 see claims 1-17 ---	1
		-/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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**INTERNATIONAL SEARCH REPORT**Int'l. Application No  
**PCT/US 93/10173****C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT**

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP,A,0 374 340 (BASF CORPORATION) 27 June 1990 see claims 1-8 -----	1
A	US,A,4 147 679 (R. L. SERIVEN) 3 April 1979 see claim 1 -----	1

## INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 93/10173

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO-A-8700851	12-00-00	CA-A-	1295767	11-02-92
		EP-A-	0233253	26-08-87
		JP-B-	1049747	25-10-89
		JP-T-	63501576	16-06-88
US-A-3984382	05-00-00	NONE		
US-A-3959201	25-00-00	US-A-	3994851	30-11-76
		CA-A-	1028079	14-03-78
		DE-A-	2447812	17-04-75
		FR-A, B	2247518	09-05-75
		GB-A-	1491014	09-11-77
		JP-C-	931912	07-11-78
		JP-A-	50066529	04-06-75
		JP-B-	52000971	11-01-77
WO-A-9108269	13-06-91	DE-A-	3940316	13-06-91
		AU-A-	6894391	26-06-91
		CA-A-	2070807	07-06-91
		EP-A-	0502934	16-09-92
EP-A-0374340	27-00-00	US-A-	4837278	06-06-89
US-A-4147679	03-00-00	NONE		